What is claimed is:

- 1. A method of producing a biotin vitamer by:
- (a) culturing a bacterium comprising a lysineutilizing DAPA aminotransferase, said culturing taking place in an environment anriched for lysine, a lysine analog, or a lysine precursor; and
 - (b) recovering said biotin vitamer.
- 2 . (a) culturing a bacterium comprising a lysine-
- 3 utilizing DAPA aminotransferase, wherein said bacterium is
- 4 deregulated with respect to lysine production; and
- (b) recovering said biotin vitamer.
- 1 3. The method of claim 1 in which the bacterium is
- 2 engineered to overproduce a lysine-utilizing DAPA
- 3 aminotransferase.

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- 1 4. The method of claim 2 in which the bacterium is
- 2 engineered to overproduce a lysine-utilizing DAPA
- 3 aminotransferase.
- 1 5. The method of claim 2 or claim 4, wherein
- 2 lysine, a lysine analog, or a lysine precursor is
- 3 exogenously added to the culture.
- 1 1 6. The method of claim 1, claim 2, claim 3, or
- 2 claim 4, in which xsine, a lysine analog, or a lysine
- 3 precursor is exogenously added to the culture and totals at
- 4 least 10 mmoles per liter of culture.

- 7. The method of claim 1, claim 2, claim 3, or claim 4, in which the biotin vitamer is biotin, dethiobiotin, or diaminopelargonic acid (DAPA).

 8. The method of claim 1, claim 2, claim 3, or claim 4, in which the biotin vitamer is dethiobiotin, and,
- claim 4, in which the biotin vitamer is dethiobiotin, and,
 after recovering the dethiobiotin, the method further
 comprises converting the recovered dethiobiotin to biotin by
 a separate fermentation, biochemical reaction, or chemical
 reaction, and recovering biotin.
- 9. The method of claim 1, claim 2, claim 3, or claim 4, in which the bacterium is resistant to a lysine analog.
- 1 10. The method of claim 9, wherein said analog is 2 S-2-aminoethyl-L-cysteine (AEC).
- 1 30 11. The method of claim 1 or claim 2, in which the bacterium is deregulated with respect to at least one biotin synthetic pathway step in addition to bioA expression.
- 1 12. The method of claim 1, claim 2, claim 3, or 2 claim 4, in which the biotin vitamer is biotin, and the 3 method comprises recovering and purifying the biotin.
- 1 3. The method of claim 1, claim 2, claim 3, or claim 4, wherein said bacterium is further engineered to produce a SAM-utilizing DAPA aminotransferase.
- 1 14. The method of claim 13 in which methionine, S-2 adenosylmethionine (SAM), or an analog of SAM is added to 3 the culture.

- 1 15. The method of claim 13 wherein lysine, a lysine
- 2 analog, or a lysine precursor is added to the culture.
- 1 16. The method of claim 14, wherein lysine, a
- 2 lysine analog, or a lysine precursor is added to the
- 3 culture.
- 1 17. The method of claim 15 in which lysine or a
- 2 lysine analog exogenously added to the culture totals at
- 3 least 10 mmoles per liter of culture.
- 1 18. The method of claim 16 in which lysine or a
- 2 lysine analog exogenously added to the culture totals at
- 3 least 10 mmoles per liter of culture.
- 1 19. The method of claim 13 in which the biotin
- 2 vitamer is biotin, dethiobiotin, or diaminopelargonic acid
- 3 (DAPA).
- 1 20. The method of claim 13 in which the biotin
- 2 vitamer is dethiobiotin, and, after recovering the
- 3 dethiobiotin, the method further comprises converting the
- 4 recovered dethiobiotin to biotin by a separate fermentation,
- 5 biochemical reaction, or chemical reaction, and recovering
- 6 biotin.
- 1 $\rightarrow h$ 21. The method of claim 13 in which the bacterium
- 2 is deregulated with respect to at least one biotin synthetic
- 3 pathway step other than bioA expression.
- 1 22. The method of claim 13 in which the biotin
- 2 vitamer is biotin, and the method comprises recovering and
- 3 purifying the biotin.

| 1 | 23. A bacterium engineered to overproduce a lysine- |
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| 2 | utilizing DAPA aminotransferase and a SAM-utilizing DAPA |
| 3 | aminotransferase. |
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| 1 | 24. The bacterial strains BI90 (ATCC) and BI96 |
| 2 | (ATCC). |
| - | (III 66, I |
| 1 | 25. The bacterium of claim 23, wherein the strain |
| 2 | is further engineered to overproduce the biotin vitamer by |
| 3 | engineered deregulation of at least one biotin synthetic |
| | step, in addition to bioA expression. |
| 4 | step, in addition to blow expression. |
| - | 26. The bacterial strain BI603 (ATCC). |
| 1 | 26. The pacternal strain broos (Arec/. |
| _ | on a best with ancincored to everproduce a lygine- |
| 1 | 27. A bacterium engineered to overproduce a lysine- |
| 2 | utilizing DAPA aminotransferse, wherein the bacterium is |
| 3 | further engineered to overproduce lysine. |
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| 1 | 28. The bacterial strain BI641 (ATCC) or BI642 |
| 2 | (ATCC). |
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| 1 | 29. A biotin vitamer manufactured by the method of |
| 2 | claim 1, claim 2, claim 3, or claim 4. |
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| 1 | 30. A biotin vitamer manufactured by the method of |
| 2 | claim 13. |
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| 1 | 31. A biotin vitamer manufactured by the method of |
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claim 14.

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